BP WASH- benzoyl peroxide emulsion CINTEX SERVICES, LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

BP Wash 10% (benzoyl peroxide 10%)

Active ingredient

Benzoyl peroxide 10%

Purpose

Acne treatment

Use

For the treatment of acne

Warnings

For external use only

Do not use if you

- have very sensitive skin
- are sensitive to benzovl peroxide

When using this product

- skin irritation and dryness is more likely to occur if you use another topical acne medication at the same time. If irritation occurs, only use one topical acne medication at a time.
- avoid unnecessary sun exposure and use a sunscreen
- avoid contact with the eyes, lips, and mouth
- avoid contact with hair and dyed fabrics, which may be bleached by this product
- skin irritation may occur, characterized by redness, burning, itching, peeling, or possibly swelling. Irritation may be reduced by using the product less frequently or in a lower concentration.

Stop use and ask a doctor if irritation becomes severe.

KEEP THIS AND ALL DRUGS OUT OF THE REACH OF CHILDREN.

If swallowed, get medical help or contact a Poison Control Center immediately.

Directions

- clean the skin thoroughly before applying this product
- cover the entire affected area with a thin layer and rinse thoroughly one to three times daily
- because excessive drying of the skin may occur, start with one application daily, then gradually increase to two or three times daily if needed or as directed by a doctor
- if bothersome dryness or peeling occurs, reduce application to once a day or every other day.
- if going outside, apply sunscreen after using this product. If irritation or sensitivity develops, stop

use of both products and ask a doctor.

Other information

Store at 20°C to 25°C (68°F to 77°F), excursions permitted between 15°C and 30°C (between 59°F and 86°F).

Inactive ingredients

carbomer, cetearyl alcohol, cocamidopropyl betaine, dimethyl isosorbide, hydrogenated castor oil, imidurea, methylparaben, mineral oil, PEG-14M, salicylic acid, sodium cocyl isethionate, sodium hydroxide, titanium dioxide, water, zea mays (corn) starch

Questions?

1-855-899-4237



NDC 24470-901-08

BP Wash

Acne Treatment

benzoy peroxide 10%

Net wt. 8 oz (227 g)



Drug Facts

Active ingredient Benzoyl peroxide 10%.

Purpose

Acne treatment

Use For the treatment of acne

Warnings

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Other information Store at 15°-30°C (59°-36°F).

Inactive ingredients carbomer, cetearyl alcohol, cocamidopropyl betaine, dimethyl isosorbide, hydrogenated castor oil, imidurea, methylparaben, mineral oil, PEG-14M, salicylic acid, sodium cocoyl isethionate, sodium hydroxide, titanium dioxide, water, zea mays (corn) starch



Questions? 1-855-899-4237

BP WASH

benzoyl peroxide emulsion

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:24470-901

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

BENZOYL PEROXIDE (UNII: W9 WZN9 A0 GM) (BENZOYL PEROXIDE - UNII: W9 WZN9 A0 GM) | BENZOYL PEROXIDE | 100 mg in 1 g

Inactive Ingredients

mactive ingredients		
Ingredient Name	Strength	
COCAMIDO PRO PYL BETAINE (UNII: 50 CF30 11KX)		
MINERAL OIL (UNII: T5L8T28FGP)		
POLYETHYLENE OXIDE 600000 (UNII: 2126FD486L)		
HYDRO GENATED CASTOR OIL (UNII: ZF94AP8MEY)		
WATER (UNII: 059QF0KO0R)		
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)		
IMIDUREA (UNII: M629807ATL)		

SODIUM COCOYL ISETHIONATE (UNII: 518 XTE8 493)	
STARCH, CORN (UNII: O8232NY3SJ)	
CARBO XYPO LYMETHYLENE (UNII: 0 A5MM307FC)	
DIMETHYL ISOSORBIDE (UNII: SA6A6V432S)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)	

Product Characteristics			
Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:24470-901-05	142 g in 1 BOTTLE; Type 0: Not a Combination Product	0 1/15/20 12	
ı	2 NDC:24470-901-08	227 g in 1 BOTTLE; Type 0: Not a Combination Product	0 1/15/20 12	

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph final	part333D	0 1/15/20 12				

Labeler - CINTEX SERVICES, LLC (078304114)

Revised: 12/2020 CINTEX SERVICES, LLC